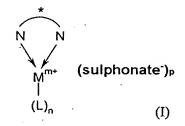
WHAT IS CLAIMED IS:

- 1. Substances comprising at least
 - one micro-, meso- or macroporous support material and
 - compounds, adsorbed thereon or therein, of the formula (I)



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where

N . N

is an enantiomerically enriched chiral nitrogen compound,

 (M^{m+})

is a metal having valency m

L

is an anionic or uncharged ligand

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(sulphonate) is the anion of a sulphonic acid and

p

is one or two and

n

is one, two, three or four,

with the proviso that m-p-[number of anionic ligands] = 0.

- Substances according to Claim 1, characterized in that the support
 materials have a pore size of 15 to 250 Å.
 - 3. Substances according to Claim 1, characterized in that the support materials are silica gels or zeolites of the MOR, X, Y, MCM, ZSM, FAU,

MFI, L, BEA, FER, A and SBA, AlPO, MAlPO or SAPO type, and the zeolites are optionally isomorphically substituted.

4. Substances according to Claim 1, characterized in that, in formula (I),

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is enantiomerically enriched chiral nitrogen compounds of the formula (II)

$$R^{2}$$
 $N-R^{3}-N$ R^{5} (II)

where

 R^1 , R^2 , R^4 and R^5 are each independently hydrogen, C_1 - C_8 -alkyl, C_5 - C_{15} -arylalkyl, C_4 - C_{14} -aryl, or NR^1R^2 and/or NR^4R^5 as a whole is a cyclic amino radical having a total of 4 to 20 carbon atoms,

R³ is a divalent radical having 2 to 30 carbon atoms or

R³ and at least one of the radicals R¹, R², R⁴ and R⁵ together are part of a cyclic amino radical having a total of 4 to 20 carbon atoms.

5. Substances according to Claim 4, characterized in that

15 R¹, R², R⁴ and R⁵ are each independently hydrogen, C₁-C₈-alkyl, C₅-C₁₅-arylalkyl or C₄-C₁₄-aryl, or NR¹R² and/or NR⁴R⁵ as a whole is a 5-or 6-membered monocyclic amino radical which is optionally mono-, di-, tri- or tetrasubstituted on the carbon framework by C₁-C₄-alkyl and

- R³ is a divalent radical which is selected from the group of C₂-C₈-alkylene which may optionally be further mono- or diubstituted by C₄-C₁₄-aryl radicals, C₅-C₁₅-arylalkylene, C₄-C₁₄-arylene or bis(C₄-C₁₄-arylene) or
- R³ and one of the radicals R¹, R², R⁴ and R⁵ together are part of a 5- or 6-membered monocyclic amino radical which is optionally additionally mono-, di-, tri- or tetrasubstituted on the carbon framework by C₁-C₄-alkyl.
- 6. Substances according to Claim 1, characterized in that, in formula (I),

 (M m+) is cobalt in the formal oxidation states 0, +2 and +3, rhodium and iridium in the formal oxidation states +1 and +3, nickel, palladium and platinum in the formal oxidation states 0 and +2 or ruthenium in the formal oxidation state +2.
- Substances according to Claim 1, characterized in that, in formula (I),
 L is the following types of ligand: monoolefins, diolefins, nitriles,
 aromatics or anionic ligands.
 - 8. Substances according to Claim 1, characterized in that (sulphonate-) is salts of the type R₆SO₃- where R⁶ is C₁-C₁₂-alkyl, C₁-C₂₀-haloalkyl, C₄-C₁₄-aryl or C₅-C₁₅-arylalkyl.
- 9. Substances according to Claim 1, characterized in that compounds of the formula (I) are those of the formulae (Ia), (Ib), (Ic), (Id), (Ie) and (If)

$$H_2N$$
 NH_2
 M^+ sulphonate

where, in each case,

- * marks a stereogenic centre which is either R- or S-configured, with the proviso that mesoforms are excluded (compounds of the formula (Ic) and (Id))
- 5 M⁺ is rhodium^I or iridium^I and
 - L is cod or nbd and

(Ie),

- sulphonate is trifluoromethanesulphonate, mesylate or nonafluorobutanesulphonate.
- 10. Compounds of the formula (I) as defined in Claim 1, with the exception of the following compounds:
- 5 [Rh(cod)((S)-2-aminomethyl-1-ethylpyrrolidine)]OTf and [Rh(cod)((1R,2R)-1,2-diphenylethylenediamine)]OTf.
 - 11. A process for conducting asymmetric reactions comprising catalyzing the reactions with substances according to Claim 1.
 - 12. Catalysts comprising substances according to Claim 1.
- 10 13. Process for preparing enantiomerically enriched compounds comprising catalyzing the preparation with catalysts according to Claim 12.
 - 14. Process according to Claim 13, characterized in that processes for catalytically preparing enantiomerically enriched compounds are asymmetric hydrogenations.
- 15 Process according to Claim 14, characterized in that asymmetric hydrogenations are hydrogenations of prochiral C=C bonds, C=O bonds and C=N bonds.
- Process according to Claim 15, characterized in that hydrogenations of prochiral C=O bonds are hydrogenations of α- and β-keto carboxylic esters.
 - 17. Process according to Claim 16, characterized in that α and β -keto carboxylic esters are those of the formula (IV)

where

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 R^6 and R^8 are each independently $C_1\text{-}C_{12}\text{-}alkyl,\,C_1\text{-}C_{12}\text{-}haloalkyl\,C_5\text{-}C_{15}\text{-}$ arylalkyl or $C_4\text{-}C_{14}\text{-}aryl$ and

 R^7 is absent or is 1,1-(C_1 - C_4 -alkylene).

- 18. Process according to Claim 14, characterized in that the reaction temperature for asymmetric hydrogenations is 0 to 200°C and the partial hydrogen pressure is 0.1 to 200 bar.
- 19. Process according to Claim 14, characterized in that the asymmetric

 hydrogenations is conducted in the presence of solvents which are aliphatic or aromatic, optionally halogenated, hydrocarbons, ethers and/or alcohols.
 - 20. Process according to Claim 13, characterized in that the weight ratio of catalysts according to Claim 1 to substrate is 1:1 to 1:10 000.